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OM protein - protein search, using sw model

Run on: October 1, 2002, 06:24:15 ; Search time 32.44 seconds
(without alignments)
1222.360 Million cell updates/sec

Title: US-09-522-752-2

Perfect score: 1854

Sequence: 1 MADDYGSSTSSMEDYVNF.....EGSLKLSMLLETSGALS 357

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 747574 seqs, 111073796 residues

Tc number of hits satisfying chosen parameters: 747574

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : A_Geneseq_032802.*

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3: /SIDSL1/gcgdata/hold-geneseg/geneseq-emb1/AA1982.DAT.*
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22: /SIDSL1/gcgdata/hold-geneseg/geneseq-emb1/AA2001.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	1854	100.0	357	21	Human CC chemokine
2	1854	100.0	357	21	Human G protein-co
3	1854	100.0	357	22	Human CCR9b protei
4	1854	100.0	369	22	Human CCR9a protei
5	1848	99.7	357	21	Human mutant G pro
6	1848	99.7	369	22	Non-endogenous hum
7	780.5	42.1	358	15	Partial sequence o
8	780.5	42.1	358	21	Human 7TM receptor
9	780.5	42.1	378	19	Human V31 seven tr
10	780.5	42.1	378	21	Human 7TM receptor
11	780.5	42.1	378	21	Human G protein-co

12	780.5	42.1	378	22	AAG80114	Human CCR7 protein
13	780.5	42.1	378	22	AAR50859	Human CCR7. Homo
14	780.5	42.1	410	15	AAR53743	Putative seven tra
15	780.5	42.1	410	19	AAW48723	Polypeptide sequen
16	780.5	42.1	410	21	ABG21687	Genomic clone of 7
17	780.5	42.1	569	22	ABG12373	Novel human diagno
18	777.5	41.9	378	15	AAR53744	Putative seven tra
19	776.5	41.9	378	21	AAV90663	Human mutant G pro
20	758.5	40.9	378	15	AAR54079	Epstein Barr virus
21	758.5	40.9	378	19	AAW56164	G-protein coupled
22	758.5	40.9	378	19	AAW53622	Epstein Barr virus
23	757.5	40.9	378	21	AAW21699	7TM receptor prote
24	742	40.0	359	15	AAR53747	Seven transmembran
25	742	40.0	359	19	AAW48728	Murine V31 seven t
26	742	40.0	359	21	AAW21691	Murine 7TM recepto
27	721	38.9	361	20	AAW97348	An Epstein-barr vi
28	671.5	36.2	351	20	AAW23825	A7 times membrane
29	652	35.2	350	22	AAW67238	Amino acid sequenc
30	640	34.5	350	20	AAV57291	Mouse BCCR protei
31	638.5	34.4	369	22	AAW80113	Human CCR6b protei
32	638.5	34.4	374	22	AAG80112	Human CCR6a protei
33	637	34.4	349	20	AAW93170	Human HFA041 prot
34	637	34.4	350	20	AAW57290	Human BCCR protei
35	637	34.4	350	20	AAW30125	A human seven-pass
36	637	34.4	350	20	AAV17435	Human signal pepti
37	637	34.4	350	20	AAW93169	Human HFA041 prot
38	637	34.4	350	21	AAV94325	Human seven transm
39	637	34.4	350	22	AAG80119	Human CCR11 protei
40	637	34.4	350	22	AAU08994	Human G protein-co
41	637	34.4	350	22	AAG67237	Amino acid sequenc
42	637	34.4	382	22	AAW62389	Human chemokine re
43	635	34.3	333	20	AAV57289	Human BCCR, partia
44	632	34.1	350	21	AAV71301	Human orphan G pro
45	632	34.1	350	21	AAW02835	Human G protein co

ALIGNMENTS

RESULT 1

AAAB19605
ID AAB19605 standard; Protein; 357 AA.
XX
XX AAB19605;
AC
XX
DT 22-JAN-2001 (first entry)
XX
DE Human CC chemokine receptor GPR-9-6.
KW GPR-9-6; human; chemokine receptor; TECK; cancer; leukaemia;
KW lymphoma; carcinoma; inflammation; Crohn's disease; colitis;
KW therapy; diagnosis.
XX
OS Homo sapiens.
XX
FN WO200053635-A1.
XX
PD 14-SEP-2000.
XX
PF 10-MAR-2000; 2000WO-US06240.
XX
PR 11-MAR-1999; 99US-0266464.
XX
PA (LEUK-) LEUKOSITE INC.
PI Andrew DP, Zabel BA, Ponath PD;
XX
DR WPI; 2000-572263/53.
XX
PT Antibody or its antigen-binding fragment which binds to the mammalian
PT CC chemokine receptor GPR-9-6, useful for treating inflammatory
PT diseases, cancer or inhibiting GPR-9-6-mediated homing of leukocytes to
PT mucosal tissue -

XX PS Disclosure; Fig 14A-B; 114pp; English.

XX CC The present sequence is that of human GPR-9-6, a CC chemokine

CC CC receptor that is expressed on the majority of thymocytes and also

CC CC on a subset of memory CD4 lymphocytes that traffic to mucosal

CC CC sites, suggesting a dual role in T cell development and mucosal

CC CC immune response. The invention relates to an antibody that binds

CC CC to GPR-9-6 and blocks the binding of a ligand, such as TECK (see

CC CC AAB19607), to the receptor. Also provided is a method of identifying

CC CC agents which can bind to GPR-9-6 and inhibit the binding of a

CC CC ligand and/or modulate a function of GPR-9-6. The antibodies can

CC CC be used to detect or measure expression of GPR-9-6 receptor. They

CC CC are useful for treating an inflammatory disease, cancer and

CC CC inhibiting GPR-9-6-mediated homing of leukocytes to mucosal tissue.

CC CC The cancer treated is acute or chronic leukaemia (e.g., acute T-cell

CC CC lymphoblastic leukaemia, acute B-cell lymphoblastic leukaemia,

CC CC chronic T-cell lymphoblastic leukaemia, chronic B-cell lymphoblastic

CC CC leukaemia), lymphoma (e.g., Hodgkin's disease, T cell lymphoma) or

CC CC carcinoma (e.g. breast, melanoma, myeloma, or adenoma). The

CC CC inflammatory diseases treated are Crohn's disease, colitis

CC CC (claimed), inflammatory bowel disease, mastitis, vaginitis,

CC CC cholangitis or pericholangitis, chronic bronchitis, asthma, graft

CC CC versus host disease, hypersensitivity pneumonitis, collagen

CC CC diseases, sarcoidosis, and other idiopathic conditions. Other

CC CC diseases that can be treated by the antibodies are autoimmune

CC CC diseases (e.g. rheumatoid arthritis, multiple sclerosis), infectious

CC CC diseases (e.g. bacterial and viral infections), atherosclerosis,

CC CC restenosis, AIDS, pancreatitis, insulin-dependent diabetes mellitus,

CC CC and diseases in which angiogenesis or neovascularization play a role.

XX CC Sequence 357 AA;

XX PS Query Match 100.0%; Score 1854; DB 21; Length 357;

XX PS Best Local Similarity 100.0%; Pred. No. 1.4e-191;

XX PS Matches 357; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 MADDYGSSTSSMEDYVNFDFYCEKNNVRQFASHFLPPLYWLVTIVGALNSLVILV 60

Db 1 maddygsstssmedyvfnfdfyceknvrgfashflpplwlvfivgalnslvllv 60

Qy 61 YWYCTRVKTMDFLLNLAIADLLFLVLPFWAIAAADQWKFQTFMCKVNSMYKMFYS 120

Db 61 ywycrtvktmdfllnlaiadllflvlpfwaiaaadqwkftcmckvnsmykmfys 120

Qy 121 CVLLIMCISVDRIYIAQAMRAHTWREKRLYSKMVCFIIWLAALCIPILYSQKEE 180

Db 121 cvllimcislvdryiaiaqamrahtwrekrllyskmvctfiwlaaalcipeillyskee 180

Qy 181 SGIAICTMVYPSDESTKLKSAVLTKLVILGFFLPFVVMACCYTIITHTLIQAKSSKHA 240

Db 181 sgiaictmvpsdestklksavtlklvlgfflpfvvmaccytiithtliqaksskhka 240

Qy 241 LKVTITVTLTVFLVQLPNCILLVQTDIYAMFISNCAVSTNIDICQVOTIAFFHSCL 300

Db 241 lkvtitvltvflvqlpncillvqtdiyamfiscavstnidicqvotiaffhscl 300

Qy 301 NPVLVYVGERFRDLVKTLKLCISQAQWVSFTTRREGSKLSMLLETTSGLSL 357

Db 301 npvlvyvgerfrdlvktlklcisaqawvsftrregsklssmllettsgalsl 357

RESULT 2

AAAY90615

ID AAAY90615 standard; Protein; 357 AA.

XX CC

AC AAAY90615;

XX CC

DT 21-AUG-2000 (first entry)

XX CC

DE Human G protein-coupled receptor GPR9-6.

XX CC

KW G protein-coupled receptor; GPCR; constitutively active; intracellular loop 3; transmembrane domain 6; drug screening; agonist; antagonist.

XX Homo sapiens.

OS WO2000022129-A1.

PN 20-APR-2000.

XX 12-OCT-1999; 99WO-US23938.

XX 13-OCT-1998; 98US-0170496.

PR (AREN-) ARENA PHARM INC.

XX Behan DP, Chalmers DT, Liaw CW;

XX WPI; 2000-329165/28.

DR N-PSDB; AAA30596.

XX Non-endogenous constitutively activated human G protein-coupled receptors, useful for identifying agonists for use as pharmaceutical agents

PT Example 1; Page 119-120; 341pp; English.

XX The invention relates to constitutively active, non-endogenous versions of endogenous human orphan G protein-coupled receptors (GPCRs, AAAY90643-AAAY90677 and AAAY90683-Y90687), and to DNA encoding them (AAAY30709-A30743 and AAAY30775-A30779). The mutant proteins of the invention contain a mutation in a portion of the protein comprising intracellular loop 3 (IC3) and transmembrane domain 6 (TM6). A non-endogenous amino acid, X, is substituted for an endogenous residue in IC3 at a position 16 amino acids N-terminal of an endogenous residue in TM6 to form a sequence X-(AA)15-Pro. The endogenous amino acid is selected from Lys, His, Arg or Ala, and is preferably Lys. When the endogenous residue at this position is Lys, this residue is replaced by His, Arg or preferably Ala. The 15 amino acid stretch between the substituted amino acid and the Pro may be endogenous, non-endogenous, or a mixture of endogenous and non-endogenous residues. The constitutively active GPCRs are useful for identifying antagonists, agonists and partial agonists for use as pharmaceutical agents. The mutant proteins are also useful in research settings for elucidating the roles of the receptors in normal and diseased conditions. Antagonists for a particular GPCR are useful for treating diseases and disorders associated with that receptor. Because the novel mutant GPCRs are constitutively active, they can be used directly for screening of compounds without the need for endogenous ligands. The present sequence represents a human wild-type GPCR referred to in an exemplification of the invention.

XX Sequence 357 AA;

XX PS Query Match 100.0%; Score 1854; DB 21; Length 357;

XX PS Best Local Similarity 100.0%; Pred. No. 1.4e-191;

XX PS Matches 357; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 MADDYGSSTSSMEDYVNFDFYCEKNNVRQFASHFLPPLYWLVTIVGALNSLVILV 60

Db 1 maddygsstssmedyvfnfdfyceknvrgfashflpplwlvfivgalnslvllv 60

Qy 61 YWYCTRVKTMDFLLNLAIADLLFLVLPFWAIAAADQWKFQTFMCKVNSMYKMFYS 120

Db 61 ywycrtvktmdfllnlaiadllflvlpfwaiaaadqwkftcmckvnsmykmfys 120

Qy 121 CVLLIMCISVDRIYIAQAMRAHTWREKRLYSKMVCFIIWLAALCIPILYSQKEE 180

Db 121 cvllimcislvdryiaiaqamrahtwrekrllyskmvctfiwlaaalcipeillyskee 180

Qy 181 SGIAICTMVYPSDESTKLKSAVLTKLVILGFFLPFVVMACCYTIITHTLIQAKSSKHA 240

Db 181 sgiaictmvpsdestklksavtlklvlgfflpfvvmaccytiithtliqaksskhka 240

QY 241 LKVTITVTLTVFVLSQFPYNCILLVQTTIDAYAMFISNCAVSTNIDICFQVTTIAFFHSL 300
 Db 241 lkvtitvltvfvlsqfpynclllvgtidayamfiscavstnidicfqvtqiaffhsl 300
 QY 301 NPVLYVFGFRDLVKTLKNLGCISQAQWVSTRREGSKLSMLETTSGLSL 357
 Db 301 npvlyvfgfrdlvktlknlgcisgaqvwstrregsklsmlettsgalsl 357

RESULT 3

AAG80117
 ID AAG80117 standard; Protein; 357 AA.

XX
 AC AAG80117;

DT 17-JAN-2002 (first entry)

DE Human CCR9b protein.

XX Chemokine; tumour diagnosis; colorectal; prostatic; organ rejection;
 KW inflammation; autoimmune disease; metastasis; bronchial asthma; lupus;
 KW chronic bowel inflammation; rheumatoid arthritis; cytostatic;
 KW antiinflammatory; antiasthmatic; immunosuppressive; dermatological;
 KW antirheumatic; antiarthritic.

OS Homo sapiens.

XX WO200172830-A2.

XX 04-OCT-2001.

XX 02-APR-2001; 2001WO-EP03708.

XX 31-MAR-2000; 2000DE-1016013.

XX (IPFP-) IPF PHARM GMBH.

XX (FORS/) FORSMANN U.

XX Forssmann W, Adermann K, Heitland A, Spodsborg N;

XX WPI; 2001-626256/72.

XX Diagnostic agent containing two or more receptor-specific ligands,
 PT useful for detecting tumors, inflammation etc., also therapeutic use of
 PT ligand inhibitors -

XX Disclosure; Page 11; 26pp; German.

XX This invention describes a novel diagnostic agent (A) comprising at least
 CC two different ligands (I) for receptors (II) that are implicated in
 CC disease. (A) are used for the diagnosis of tumors (especially colorectal
 CC or prostatic), organ rejection, inflammation and autoimmune diseases.
 CC Also inhibitors of (I) are used therapeutically against tumors (and their
 CC metastases), inflammation (particularly bronchial asthma or chronic bowel
 CC inflammation), or autoimmune diseases (rheumatoid arthritis or lupus),
 CC where the (cardio)vascular, lymphatic, respiratory, nervous, digestive,
 CC endocrine, motor or urogenital systems or skin are affected, and bone
 CC marrow diseases. The products of the invention are chemokine derivatives
 CC which have cytostatic, antiinflammatory, antiasthmatic,
 CC immunosuppressive, dermatological, antirheumatic, antiarthritic.
 CC Chemokines act on specific tumor and inflammatory cells through a
 CC constellation of chemokine receptors (CR), which control migration and
 CC proliferation of these cells. AAG80045-AAG80128 represent human chemokine
 CC fragments used to illustrate the method of the invention.

XX Sequence 357 AA;

Query Match 100.0%; Score 1854; DB 22; Length 357;
 Best Local Similarity 100.0%; Pred. No. 1.4e-191;
 Matches 357; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MADDYGSESTSSMEDYVNFNFTDFYCEKNVVRQFASHFLPPLYWLVFIVGALGNSLVLY 60
 Db 1 maddygsestssmedyvnfnftdfyceknvrqfashflpplywlvfivgalgnslvly 60
 QY 61 YWYTRVKTWTDMLLNLALADLLFLVLPFWAIAAADQWKFOFMCKVNVSMYKMFYS 120
 Db 61 ywytrvktwtdmllnlaladllflvlpfwaiaaadqwkfqfmckvvnsmymkmyfs 120
 QY 121 CVLLIMCISVDVRYIAIAQAMRAHTWRKRLYSKMWCFITWLAALCIPILYSQIKEE 180
 Db 121 cvllimcisdryiaiaqamrahtwrekrilyskmvcftiwlalalcipellysqikee 180
 QY 181 SGIAICTWYPSDESTKLKSAVLTLKVLGFLPFPVVMACCYTIITLQAKKSSKHKA 240
 Db 181 sgiaictmypsdestklksavtlkvlglfplfpvfmaccytiitlclqakskshka 240
 QY 241 LKVTITVTLTVFVLSQFPYNCILLVQTTIDAYAMFISNCAVSTNIDICFQVTTIAFFHSL 300
 Db 241 lkvtitvltvfvlsqfpynclllvgtidayamfiscavstnidicfqvtqiaffhsl 300
 QY 301 NPVLYVFGFRDLVKTLKNLGCISQAQWVSTRREGSKLSMLETTSGLSL 357
 Db 301 npvlyvfgfrdlvktlknlgcisgaqvwstrregsklsmlettsgalsl 357

RESULT 4

AAG80116
 ID AAG80116 standard; Protein; 369 AA.

XX
 AC AAG80116;

DT 17-JAN-2002 (first entry)

XX Human CCR9a protein.

XX Chemokine; tumour diagnosis; colorectal; prostatic; organ rejection;
 KW inflammation; autoimmune disease; metastasis; bronchial asthma; lupus;
 KW chronic bowel inflammation; rheumatoid arthritis; cytostatic;
 KW antiinflammatory; antiasthmatic; immunosuppressive; dermatological;
 KW antirheumatic; antiarthritic.

OS Homo sapiens.

XX WO200172830-A2.

XX 04-OCT-2001.

XX 02-APR-2001; 2001WO-EP03708.

XX 31-MAR-2000; 2000DE-1016013.

XX (IPFP-) IPF PHARM GMBH.

XX (FORS/) FORSMANN U.

XX Forssmann W, Adermann K, Heitland A, Spodsborg N;

XX WPI; 2001-626256/72.

XX Diagnostic agent containing two or more receptor-specific ligands,
 PT useful for detecting tumors, inflammation etc., also therapeutic use of
 PT ligand inhibitors -

XX Disclosure; Page 11; 26pp; German.

XX This invention describes a novel diagnostic agent (A) comprising at least
 CC two different ligands (I) for receptors (II) that are implicated in
 CC disease. (A) are used for the diagnosis of tumors (especially colorectal
 CC or prostatic), organ rejection, inflammation and autoimmune diseases.
 CC Also inhibitors of (I) are used therapeutically against tumors (and their
 CC metastases), inflammation (particularly bronchial asthma or chronic bowel
 CC inflammation), or autoimmune diseases (rheumatoid arthritis or lupus),
 CC where the (cardio)vascular, lymphatic, respiratory, nervous, digestive,
 CC endocrine, motor or urogenital systems or skin are affected, and bone

CC marrow diseases. The products of the invention are chemokine derivatives
CC which have cytostatic, antiinflammatory, antiasthmatic,
CC immunosuppressive, dermatological, antirheumatic, antiarthritic.
CC Chemokines act on specific tumor and inflammatory cells through a
CC constellation of chemokine receptors (CR), which control migration and
CC proliferation of these cells. AAG80045-AAG80128 represent human chemokine
CC fragments used to illustrate the method of the invention.
XX
SQ Sequence 369 AA;

Query Match 100.0%; Score 1854; DB 22; Length 369;
Best Local Similarity 100.0%; Pred. No. 1.4e-191;
Matches 357; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MADDYGSSTSMEDYVNFNFTDFCEKNVQFASHFLPPLYWLVFVIGALNSLVILV 60
DB 13 maddygsestsmedyvnfntdfyceknvrfqashflppllywlvfivgalsnlv 72
61 YWYCTRVKTMDFLLNLAIADLLFLVLPFWAIAAADQWKFTQFMCKVNSMYKMFYS 120
73 ywycrvktmtdmflnlaiadllflvlpfwaiaaadqwkftfmcvkvnsmykmfys 132
QY 121 CVLLIMCISVDRIYIAQAQAMRAHTWREKRLLYSKMVCFTIWLAAALCIPILYSQIKEE 180
DB 133 cvllimcisdryiaiaqamrahtwrekrllyskmvcftiwlaaalcipellysqikee 192
QY 181 SGIAICTMVYPSPDESTKLKSAVLTLLKVLGFPFVVMACCVTIIHTLIQAKSSKHKA 240
DB 193 sgiaictmvypsdetkksavltllkvlgfflvfvmvaccvtiilhtliqaksskhka 252
QY 241 LKVTITVTLTVFVLSQFPYNCILLVQTIDAYAMFISNCAVSTNIDICFQVTTOTIAFFHSCL 300
DB 253 lkvtitvltvflvsqfpyncillvqtidayamfiscavstnidicfvtqtiaffhscl 312
QY 301 NPVLYVFGFRFRDLVKTLLKGLCISQAQWVSFTRREGSLKLSMLLETTSGLSL 357
DB 313 npvlyvfgfrfrdlvktllkglcisaqawvsftrregsklssmllettsgalsl 369

RESULT 5
AAY90649
ID AAY90649 standard; Protein; 357 AA.
XX
AC AAY90649;
XX
DT 21-AUG-2000 (first entry)
XX
Human mutant G protein-coupled receptor GPR9-6 (L241K).
G protein-coupled receptor; GPCR; constitutively active;
intracellular loop 3; transmembrane domain 6; drug screening;
agonist; antagonist; mutant; mutein.
XX
OS Homo sapiens.
OS Synthetic.
XX
PN WO200022129-A1.
XX
PD 20-APR-2000.
XX
PF 12-OCT-1999; 99WO-US23938.
XX
PR 13-OCT-1998; 98US-0170496.
XX
PA (AREN-) ARENA PHARM INC.
XX
PI Behan DP, Chalmers DT, Liaw CW;
XX
DR WPI; 2000-329165/28.
DR N-PSDB; AAA30715.
XX
PT Non-endogenous constitutively activated human G protein-coupled

PT receptors, useful for identifying agonists for use as pharmaceutical
PT agents
XX
PS Example 2; Page 226-227; 34lpp; English.
XX

The invention relates to constitutively active, non-endogenous versions
of endogenous human orphan G protein-coupled receptors (GPCRs, AAY90643-
AAY90677 and AAY90683-Y90687), and to DNA encoding them (AAA30709-A30743
and AAA30775-A30779). The mutant proteins of the invention contain a
mutation in a portion of the protein comprising intracellular loop 3
(IC3) and transmembrane domain 6 (TM6). A non-endogenous amino acid, X,
is substituted for an endogenous residue in IC3 at a position 16 amino
acids N-terminal of an endogenous proline in TM6 to form a sequence
X-(AA)15-Pro. The endogenous amino acid is selected from Lys, His, Arg
or Ala, and is preferably Lys. When the endogenous residue at this
position is Lys, this residue is replaced by His, Arg or preferably Ala.
The 15 amino acid stretch between the substituted amino acid and the pro
may be endogenous, non-endogenous, or a mixture of endogenous and
non-endogenous residues. The constitutively active GPCRs are useful for
identifying antagonists, agonists and partial agonists for use as
pharmaceutical agents. The mutant proteins are also useful in research
settings for elucidating the roles of the receptors in normal and
diseased conditions. Antagonists for a particular GPCR are useful for
treating diseases and disorders associated with that receptor. Because
the novel mutant GPCRs are constitutively active, they can be used
directly for screening of compounds without the need for endogenous
ligands. Sequences AAY90643- AAY90677 and AAY90683-Y90687 the mutant
human GPCRs of the invention.

XX Sequence 357 AA;

Query Match 99.7%; Score 1848; DB 21; Length 357;
Best Local Similarity 99.7%; Pred. No. 6.2e-191;
Matches 356; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 MADDYGSSTSMEDYVNFNFTDFCEKNVQFASHFLPPLYWLVFVIGALNSLVILV 60
DB 1 maddygsestsmedyvnfntdfyceknvrfqashflppllywlvfivgalsnlv 60
QY 61 YWYCTRVKTMDFLLNLAIADLLFLVLPFWAIAAADQWKFTQFMCKVNSMYKMFYS 120
DB 61 ywycrvktmtdmflnlaiadllflvlpfwaiaaadqwkftfmcvkvnsmykmfys 120
QY 121 CVLLIMCISVDRIYIAQAQAMRAHTWREKRLLYSKMVCFTIWLAAALCIPILYSQIKEE 180
DB 121 cvllimcisdryiaiaqamrahtwrekrllyskmvcftiwlaaalcipellysqikee 180
QY 181 SGIAICTMVYPSPDESTKLKSAVLTLLKVLGFPFVVMACCVTIIHTLIQAKSSKHKA 240
DB 181 sgiaictmvypsdetkksavltllkvlgfflvfvmvaccvtiilhtliqaksskhka 240
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DB 241 lkvtitvltvflvsqfpyncillvqtidayamfiscavstnidicfvtqtiaffhscl 300
QY 301 NPVLYVFGFRFRDLVKTLLKGLCISQAQWVSFTRREGSLKLSMLLETTSGLSL 357
DB 301 npvlyvfgfrfrdlvktllkglcisaqawvsftrregsklssmllettsgalsl 357

RESULT 6
ABB56344
ID ABB56344 standard; Protein; 369 AA.
XX
AC ABB56344;
XX
DT 18-FEB-2002 (first entry)
XX
Non-endogenous human GPCR protein, SEQ ID NO: 481.
DE Human; G protein-coupled receptor; GPCR; non-endogenous; mutant;
XX constitutively activated GPCR; agonist; disease.
KW

CC acids N-terminal of an endogenous proline in TM6 to form a sequence
CC X-(AA)15-Pro. The endogenous amino acid is selected from Lys, His, Arg
CC or Ala, and is preferably Lys. When the endogenous residue at this
CC position is Lys, this residue is replaced by His, Arg or preferably Ala.
CC The 15 amino acid stretch between the substituted amino acid and the Pro
CC may be endogenous, non-endogenous, or a mixture of endogenous and
CC non-endogenous residues. The constitutively active GPCRs are useful for
CC identifying antagonists, agonists and partial agonists for use as
CC pharmaceutical agents. The mutant proteins are also useful in research
CC settings for elucidating the roles of the receptors in normal and
CC diseased conditions. Antagonists for a particular GPCR are useful for
CC treating diseases and disorders associated with that receptor. Because
CC the novel mutant GPCRs are constitutively active, they can be used
CC directly for screening of compounds without the need for endogenous
CC ligands. The present sequence represents a human wild-type GPCR referred
CC to in an exemplification of the invention.

XX Sequence 378 AA;

Query Match 42.1%; Score 780.5; DB 21; Length 378;
Best Local Similarity 42.9%; Pred. No. 1.2e-75;
Matches 153; Conservative 77; Mismatches 104; Indels 23; Gaps 6;

QY 1 MADDYGESESSMEDYVNFNTDFCEKNNVRQFASHPLPLPLVWLVFVIGALGNSLVILV 60
DB 28 vtdyigdnnt-----vdytlfesalcakdvrfkwlplmysislfcvlgnglvlt 82
QY 61 YWYTRVKTMTDFLNLAIADLLVLPFWATAADQMKFQTFMCKVNSMYKMFYS 120
DB 83 yiyfkrktmtdtyllnlavadiifltpfwaysaakswvfgvhfcklifaikmsffs 142
QY 121 CVLLIMCISVDRIYIAQAAMRAHTWREKRLYSKMVCFITWVLAALCIPILYSQIKEE 180
DB 143 gmlilicisidryvaivqavsahrhrarvlliskscvgilvatvisipellysdlqrs 202
QY 181 SG--IAICTWVPYDSESTKLKSAVLTUKV---ILGFFLPFVVMACCVTTIIHTLIQAKS 235
DB 203 sseqamrcsli-----tehveafitigvaqmvigfvlplamsfcylviirtllqarnf 256
QY 236 SKHKALKVTITVLTVFVLSQPPYNCILLVQTIDAYAMFISNCAVSTNIDICFOVTOTIAF 295
DB 257 ernkaikviiavvvfvlpyngvvlactvanfnitsstcelskqlnlaydvtyslac 316
QY 296 FHSLCNPLVLYVFGFRPRDLVTKLNLGICISQ---AQWVSFTRREGSLKLSMLLE 349
DB 317 vrccvnpflyafgkvkfrndlkfklgclsgqlrqsccrh-----irssmsve 369

QY T 12

AA80114
ID AAG80114 standard; Protein; 378 AA.

XX

AC AAG80114;

XX

DT 17-JAN-2002 (first entry)

XX

DE Human CCR7 protein.

XX Chemokine; tumour diagnosis; colorectal; prostatic; organ rejection;
XX inflammation; autoimmune disease; metastasis; bronchial asthma; lupus;
XX chronic bowel inflammation; rheumatoid arthritis; cytostatic;
XX antiinflammatory; antiasthmatic; immunosuppressive; dermatological;
XX antirheumatic; antiarthritic.

OS Homo sapiens.

XX

PN WO200172830-A2.

XX

PD 04-OCT-2001.

XX

PF 02-APR-2001; 2001WO-EP03708.

XX

PR 31-MAR-2000; 2000DE-1016013.

XX

PA (IPFP-) IPF PHARM GMBH.

PA (TORS/) FORSSMANN U.

XX

PI Forssmann W, Adermann K, Heitland A, Spodsberg N;

XX

DR WPI; 2001-626256/72.

XX

PT Diagnostic agent containing two or more receptor-specific ligands,
PT useful for detecting tumors, inflammation etc., also therapeutic use of
PT ligand inhibitors

XX

PS Disclosure; Page 10; 26pp; German.

XX

CC This invention describes a novel diagnostic agent (A) comprising at least
CC two different ligands (I) for receptors (II) that are implicated in
CC disease. (A) are used for the diagnosis of tumors (especially colorectal
CC or prostatic), organ rejection, inflammation and autoimmune diseases.
CC Also inhibitors of (I) are used therapeutically against tumors (and their
CC metastases), inflammation (particularly bronchial asthma or chronic bowel
CC inflammation), or autoimmune diseases (rheumatoid arthritis or lupus),
CC where the (cardio)vascular, lymphatic, respiratory, nervous, digestive,
CC endocrine, motor or urogenital systems or skin are affected, and bone
CC marrow diseases. The products of the invention are chemokine derivatives
CC which have cytostatic, antiinflammatory, antirheumatic, antiarthritic,
CC immunosuppressive, dermatological, antitumor, antiasthmatic,
CC Chemokines act on specific tumor and inflammatory cells through a
CC constellation of chemokine receptors (CR) which control migration and
CC proliferation of these cells. AAG80045-AAG80128 represent human chemokine
CC fragments used to illustrate the method of the invention.

XX Sequence 378 AA;

Query Match 42.1%; Score 780.5; DB 22; Length 378;

Best Local Similarity 42.9%; Pred. No. 1.2e-75;

Matches 153; Conservative 77; Mismatches 104; Indels 23; Gaps 6;

QY 1 MADDYGESESSMEDYVNFNTDFCEKNNVRQFASHPLPLPLVWLVFVIGALGNSLVILV 60
DB 28 vtdyigdnnt-----vdytlfesalcakdvrfkwlplmysislfcvlgnglvlt 82
QY 61 YWYTRVKTMTDFLNLAIADLLVLPFWATAADQMKFQTFMCKVNSMYKMFYS 120
DB 83 yiyfkrktmtdtyllnlavadiifltpfwaysaakswvfgvhfcklifaikmsffs 142
QY 121 CVLLIMCISVDRIYIAQAAMRAHTWREKRLYSKMVCFITWVLAALCIPILYSQIKEE 180
DB 143 gmlilicisidryvaivqavsahrhrarvlliskscvgilvatvisipellysdlqrs 202
QY 181 SG--IAICTWVPYDSESTKLKSAVLTUKV---ILGFFLPFVVMACCVTTIIHTLIQAKS 235
DB 203 sseqamrcsli-----tehveafitigvaqmvigfvlplamsfcylviirtllqarnf 256
QY 236 SKHKALKVTITVLTVFVLSQPPYNCILLVQTIDAYAMFISNCAVSTNIDICFOVTOTIAF 295
DB 257 ernkaikviiavvvfvlpyngvvlactvanfnitsstcelskqlnlaydvtyslac 316
QY 296 FHSLCNPLVLYVFGFRPRDLVTKLNLGICISQ---AQWVSFTRREGSLKLSMLLE 349
DB 317 vrccvnpflyafgkvkfrndlkfklgclsgqlrqsccrh-----irssmsve 369

RESULT 13

AA805859

ID AAB50859 standard; protein; 378 AA.

XX

AC AAB50859;

XX

DT 16-MAR-2001 (first entry)

XX

DE Human CCR7.

GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM protein - protein search, using sw model

Run on: October 1, 2002, 06:26:10 ; Search time 12.97 Seconds
(without alignments)
672.316 Million cell updates/sec

Title: US-09-522-752-2
Perfect score: 1854
Sequence: 1 MADDYGESTSSMEDYVNFN.....EGSLKLSMLLETTSGLSL 357

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 231628 seqs, 24425594 residues

Total number of hits satisfying chosen parameters: 231628

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

- Database :
- 1: /cgn2.6/ptodata/2/1aa/5A.COMB.pap.*
 - 2: /cgn2.6/ptodata/2/1aa/5B.COMB.pap.*
 - 3: /cgn2.6/ptodata/2/1aa/6A.COMB.pap.*
 - 4: /cgn2.6/ptodata/2/1aa/6B.COMB.pap.*
 - 5: /cgn2.6/ptodata/2/1aa/PCRTUS.COMB.pap.*
 - 6: /cgn2.6/ptodata/2/1aa/backfiles1.pap.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	1854	100.0	357	4	US-09-266-464-2
2	780.5	42.1	358	1	US-08-153-848-19
3	780.5	42.1	358	3	US-09-299-843A-19
4	780.5	42.1	358	4	US-09-088-337B-19
5	780.5	42.1	358	5	PCR-US93-11153-19
6	780.5	42.1	378	1	US-08-153-848-15
7	780.5	42.1	378	3	US-09-299-843A-15
8	780.5	42.1	378	4	US-09-251-545-1
9	780.5	42.1	378	4	US-09-088-337B-15
10	780.5	42.1	378	5	PCR-US93-11153-15
11	780.5	42.1	410	1	US-08-153-848-7
12	780.5	42.1	410	3	US-09-299-843A-7
13	780.5	42.1	410	4	US-09-088-337B-7
14	780.5	42.1	410	5	PCR-US93-11153-7
15	758.5	40.9	378	1	US-08-383-750-2
16	758.5	40.9	378	1	US-08-383-751A-2
17	758.5	40.9	378	3	US-08-352-678-2
18	758.5	40.9	378	4	US-09-045-583-49
19	758.5	40.9	378	5	PCR-US93-09636-2
20	757.5	40.9	378	3	US-09-299-843A-66
21	757.5	40.9	378	4	US-09-088-337B-66
22	742	40.0	359	1	US-08-153-848-24
23	742	40.0	359	3	US-09-299-843A-24
24	742	40.0	359	4	US-09-088-337B-24
25	742	40.0	359	5	PCR-US93-11153-24
26	721	38.9	361	2	US-08-902-294-2
27	721	38.9	361	3	US-09-178-637-2

28	652	35.2	350	2	US-08-966-316-18	Sequence 18, Appl
29	638.5	34.4	374	4	US-09-045-583-48	Sequence 48, Appl
30	637	34.4	350	2	US-08-966-316-16	Sequence 16, Appl
31	624	33.7	342	4	US-09-116-498-6	Sequence 6, Appl
32	619	33.4	342	4	US-09-116-498-4	Sequence 4, Appl
33	612	33.0	342	2	US-08-742-011-2	Sequence 2, Appl
34	612	33.0	342	4	US-09-275-384B-5	Sequence 5, Appl
35	612	33.0	342	4	US-09-116-498-2	Sequence 2, Appl
36	612	33.0	342	4	US-09-449-437A-2	Sequence 2, Appl
37	598	32.3	352	4	US-09-045-583-52	Sequence 52, Appl
38	596	32.1	352	4	US-09-087-232A-13	Sequence 13, Appl
39	596	32.1	352	4	US-08-861-105-14	Sequence 14, Appl
40	596	32.1	352	4	US-08-575-967A-2	Sequence 2, Appl
41	590	31.8	352	3	US-08-466-343D-2	Sequence 2, Appl
42	585.5	31.6	360	4	US-08-875-573-20	Sequence 20, Appl
43	585.5	31.6	360	4	US-09-232-878-2	Sequence 2, Appl
44	585.5	31.6	360	4	US-09-045-583-55	Sequence 55, Appl
45	582	31.4	360	1	US-08-202-056-7	Sequence 7, Appl

ALIGNMENTS

RESULT 1
US-09-266-464-2
; Sequence 2, Application US/09266464
; GENERAL INFORMATION:
; APPLICANT: Andrew, David P.
; APPLICANT: Zabel, Brian A.
; APPLICANT: Ponath, Paul D.
; TITLE OF INVENTION: ANTI-GPR-9-6 ANTIBODIES AND METHODS OF IDENTIFYING AGENTS WHICH MODULATE GPR-9-6 FUNCTION
; FILE REFERENCE: LKS98-16
; CURRENT APPLICATION NUMBER: US/09/266.464
; CURRENT FILING DATE: 1999-03-11
; NUMBER OF SEQ ID NOS: 7
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 2
; LENGTH: 357
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-266-464-2

Query Match	100.0%;	Score	1854;	DB	4;	Length	357;
Best Local Similarity	100.0%;	Pred. No.	7.8e-162;				
Matches	357;	Conservative	0;	Mismatches	0;	Indels	0;
Qy	1	MADDYGESTSSMEDYVNFNFTDFYCEKNVROFASHFLPPLYWLVLVFGALGNSLVILV	60				
Db	1	MADDYGESTSSMEDYVNFNFTDFYCEKNVROFASHFLPPLYWLVLVFGALGNSLVILV	60				
Qy	61	YWYCTRVKTMDFLLNLAIDLFLVLPFWAIAAADOWKQFTFCMKVNSMYKMFYS	120				
Db	61	YWYCTRVKTMDFLLNLAIDLFLVLPFWAIAAADOWKQFTFCMKVNSMYKMFYS	120				
Qy	121	CVLLIMCISVDRIYIAQAMRAHTWREKRLLYSKWCVTIWVLAALCIPEILYSQIKEE	180				
Db	121	CVLLIMCISVDRIYIAQAMRAHTWREKRLLYSKWCVTIWVLAALCIPEILYSQIKEE	180				
Qy	181	SGIAICTWYPSDESTKLKSAVTLKVLGFFLPFWMACCVTIHTLIQAKSKSKHA	240				
Db	181	SGIAICTWYPSDESTKLKSAVTLKVLGFFLPFWMACCVTIHTLIQAKSKSKHA	240				
Qy	241	LKVTITVLTVFVLSOPFYNCILLVQTIDAYAMFISNCAVSTNIDICFQVQTQIAFFHSL	300				
Db	241	LKVTITVLTVFVLSOPFYNCILLVQTIDAYAMFISNCAVSTNIDICFQVQTQIAFFHSL	300				
Qy	301	NPVLVYFVGERRDLVKTLKNLGCIQAOQVSTFRREGSLKLSMLLETTSGLSL	357				
Db	301	NPVLVYFVGERRDLVKTLKNLGCIQAOQVSTFRREGSLKLSMLLETTSGLSL	357				

RESULT 2
US-08-153-848-19
; Sequence 19, Application US/08153848
; Patent No. 5759804
; GENERAL INFORMATION:
; APPLICANT: Godiska, Ronald
; APPLICANT: Gray, Patrick W.
; APPLICANT: Schweikart, Vicki L.
; TITLE OF INVENTION: No. 5759804el Seven Transmembrane Receptors
; NUMBER OF SEQUENCES: 64
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Marshall, O'Toole, Gerstein, Murray &
; STREET: 6300 Sears Tower, 233 South Wacker Drive
; CITY: Chicago
; STATE: Illinois
; COUNTRY: USA
; ZIP: 60606
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; FILING DATE: 17-NOV-1992
; APPLICATION NUMBER: US/08/153,848
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/977,452
; FILING DATE: 17-NOV-1992
; ATTORNEY/AGENT INFORMATION:
; NAME: No. 5759804and, Greta E.
; REGISTRATION NUMBER: 35,302
; REFERENCE/DOCKET NUMBER: 31794
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (312) 474-6300
; TELEFAX: (312) 474-0448
; TELEX: 25-3856
; INFORMATION FOR SEQ ID NO: 19:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 358 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-153-848-19

Query Match 42.1%; Score 780.5; DB 1; Length 358;
Best Local Similarity 42.9%; Pred. No. 8.8e-64;
Matches 153; Conservative 77; Mismatches 104; Indels 23; Gaps 6;

Qy 1 MADDYGSSTSMEDYVNFNFTDFYCEKNNVRQFASHPFLPPLYWLVFIVGALGNSLVLY 60
Db 8 VTDDYIGNDTT-----VDYTLFESLCSKDKVRNFKANFLPIMYSIICFVGLLGNLVLT 62

Qy 61 YWYCTRVKTMDFMFLNLAADLFLVLPFWATAADQWKFQTFMCKVNSMYKMFYS 120
Db 63 YIYFKRLKTMDFMFLNLAADLFLVLPFWATAADQWKFQTFMCKVNSMYKMFYS 122

Qy 121 CVLLIMCISVDRYIAIAQAMRAHTWRKRLLYSKMVCFITWVLAALCIPILYSQIKEE 180
Db 123 GMLLLCISIDRYVAIVQAVSAHRHARVLLISKLCVGIWILATVLSIPILYSQILRS 182

Qy 181 SG--IAICTWVYPSDESTKLSAVLTAKV---ILGFPLFPVVMACCVYIIHTLIQAKS 235
Db 183 SSEQAMRSLI-----TEHVEAFITIOVAQWVIGFLVPLAMFCYLIIVITLQARNF 236

Qy 236 SKHKALKVITIVLVFVLSQFPYNCILVQTDAYAMPISNCVSTNEDICFQVQTIAF 295
Db 237 ERNKAIRVIIAAVWVFIQVLPYNGVLAQTVANFNITSSTCELSKQLNADYVTSIAC 296

Qy 296 FHSCLNPVLYVVGFRFRDLVKTILKNGCISQ---AQWVSFTREGSLKSSMLLE 349
Db 298 FHSCLNPVLYVVGFRFRDLVKTILKNGCISQ---AQWVSFTREGSLKSSMLLE 349

Db 297 VRCVNPFLYAFIGVKFRNDLFLKFLKDLGCLSQBQLRWSSCRH----IRRSSMSVE 349

RESULT 3
US-09-299-843A-19
; Sequence 19, Application US/09299843A
; Patent No. 6107475
; GENERAL INFORMATION:
; APPLICANT: Godiska, Ronald
; APPLICANT: Gray, Patrick W.
; APPLICANT: Schweikart, Vicki L.
; TITLE OF INVENTION: No. 6107475el Seven Transmembrane Receptors
; NUMBER OF SEQUENCES: 66
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Marshall, O'Toole, Gerstein, Murray &
; STREET: 6300 Sears Tower, 233 South Wacker Drive
; CITY: Chicago
; STATE: Illinois
; COUNTRY: USA
; ZIP: 60606
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/299,843A
; FILING DATE:
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 09/088,337
; FILING DATE: 01-JUN-1998
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/153,848
; FILING DATE: 17-NOV-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/977,452
; FILING DATE: 17-NOV-1992
; ATTORNEY/AGENT INFORMATION:
; NAME: Jill E. Uhl
; REGISTRATION NUMBER: 43,213
; REFERENCE/DOCKET NUMBER: 27866/32059B
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (312) 474-6300
; TELEFAX: (312) 474-0448
; TELEX:
; INFORMATION FOR SEQ ID NO: 19:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 358 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-09-299-843A-19

Query Match 42.1%; Score 780.5; DB 3; Length 358;
Best Local Similarity 42.9%; Pred. No. 8.8e-64;
Matches 153; Conservative 77; Mismatches 104; Indels 23; Gaps 6;

Qy 1 MADDYGSSTSMEDYVNFNFTDFYCEKNNVRQFASHPFLPPLYWLVFIVGALGNSLVLY 60
Db 8 VTDDYIGNDTT-----VDYTLFESLCSKDKVRNFKANFLPIMYSIICFVGLLGNLVLT 62

Qy 61 YWYCTRVKTMDFMFLNLAADLFLVLPFWATAADQWKFQTFMCKVNSMYKMFYS 120
Db 63 YIYFKRLKTMDFMFLNLAADLFLVLPFWATAADQWKFQTFMCKVNSMYKMFYS 122

Qy 121 CVLLIMCISVDRYIAIAQAMRAHTWRKRLLYSKMVCFITWVLAALCIPILYSQIKEE 180
Db 123 GMLLLCISIDRYVAIVQAVSAHRHARVLLISKLCVGIWILATVLSIPILYSQILRS 182

Qy 181 SG--IAICTWVYPSDESTKLSAVLTAKV---ILGFPLFPVVMACCVYIIHTLIQAKS 235

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: October 1, 2002, 06:26:30 ; Search time 46.91 Seconds
(without alignments)
731.270 Million cell updates/sec

Title: US-09-522-752-2
Perfect score: 1854
Sequence: 1 MADDYGSESTSMEDYVNFN.....EGSLKLSMLETTSGLSL 357

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 283138 seqs, 96089334 residues

To: Number of hits satisfying chosen parameters: 283138

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : PIR_71.*
1: pir1.*
2: pir2.*
3: pir3.*
4: pir4.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	780.5	42.1	378	2 B55735	lymphocyte-specific
2	758.5	40.9	378	2 A45680	G protein-coupled
3	757.5	40.9	378	2 A55735	G protein-coupled
4	652	35.2	350	2 JN0621	G protein-coupled
5	638.5	34.4	369	2 JC5068	G protein-coupled
6	596	32.1	352	2 A43113	chemokine (C-C) re
7	585.5	31.6	360	2 A57160	chemokine (C-C) re
8	582	31.4	360	2 A53611	interleukin-8 rece
9	577	31.1	360	2 JC4587	chemokine (C-C) re
10	575	31.0	360	2 JC2443	chemokine (C-C) re
11	571	30.8	355	2 A45177	chemokine (C-C) re
12	569	30.7	374	2 I38450	chemokine (C-C) re
13	569	30.7	383	2 S55594	G protein-coupled
14	566	30.5	358	2 A53752	interleukin-8 rece
15	565.5	30.5	353	2 S28787	neuropeptide Y/pep
16	560	30.2	355	2 G02436	chemokine (C-C) re
17	558.5	30.1	350	2 A39445	interleukin-8 rece
18	558.5	30.1	355	2 J01231	interleukin-8 rece
19	557.5	30.1	352	2 A45747	neuropeptide Y/pep
20	555	29.9	359	2 I49341	MIP-1 alpha recept
21	547.5	29.5	352	2 G00048	fusin (LESTRA) - c
22	545	29.4	359	2 A48921	interleukin-8 rece
23	543.5	29.3	356	2 S42096	interleukin-8 rece
24	528	28.5	374	2 S32785	G protein-coupled
25	527.5	28.5	355	2 I49339	macrophage inflam
26	526.5	28.4	374	2 S42628	G protein-coupled
27	524	28.3	327	2 S56162	MBP15 protein - h
28	521.5	28.1	359	2 S15403	angiotensin II rec
29	521.5	28.1	372	2 S26667	G protein-coupled

30	517.5	27.9	359	2 S44425	angiotensin II rec
31	516.5	27.9	359	2 JC1104	angiotensin II rec
32	512	27.6	355	2 JC3067	G protein-coupled
33	512	27.6	367	2 JE0349	interferon-inducib
34	508.5	27.4	359	2 A48857	angiotensin II rec
35	505.5	27.3	359	2 A42656	angiotensin II rec
36	504.5	27.2	359	2 I39418	angiotensin II rec
37	504.5	27.2	359	2 JC2134	angiotensin II rec
38	501.5	27.0	359	2 JC1194	angiotensin II rec
39	500	27.0	356	2 I49340	MIP-1 alpha recept
40	498.5	26.9	359	2 JH0621	angiotensin II rec
41	494.5	26.7	359	2 JQ1516	angiotensin II rec
42	491.5	26.5	362	2 JN0694	interleukin-8 rece
43	483	26.1	354	2 A23669	orphan G protein-c
44	482.5	26.0	355	2 JC4304	G protein-coupled
45	479	25.8	354	2 B55733	G protein-coupled

ALIGNMENTS

RESULT 1

B55735
lymphocyte-specific G protein-coupled receptor EB11 - human
N;Alternate names: Burkitt's lymphoma receptor 2; Epstein-Barr virus induced protein
C;Species: Homo sapiens (man)
C;Date: 07-Jul-1995 #sequence_revision 07-Jul-1995 #text_change 19-May-2000
R;Accession: B55735; S52443
R;Schweickart, V.L.; Raport, C.J.; Godiska, R.; Byers, M.G.; Eddy Jr., R.L.; Shows, T
Genomics 23, 643-650, 1994
A;Title: Cloning of human and mouse EB11, a lymphoid-specific G-protein-coupled recep
A;Reference number: A55735; MUID:95154835
A;Accession: B55735
A;Status: preliminary
A;Molecule type: mRNA
A;Residues: 1-378 <SCH>
A;Cross-references: GB:I31581; NID:g468319; PIDN:AAA74231.1; PID:g468320
R;Burgstahler, R.; Kempkes, B.; Staube, K.; Lipp, M.
submitted to the EMBL Data Library, February 1995
A;Description: The expression of the chemokine receptor BLR2/EB11 is specifically tra
A;Reference number: S52443
A;Accession: S52443
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 21-378 <BUR>
A;Cross-references: EMBL:X84702
C;Genetics:
A;Gene: GDB:CMKBR7; EB11; BLR2; CCR7
A;Cross-references: GDB:342065; OMIM:600242
A;Map position: 17q12-17q21.2
C;Superfamily: vertebrate rhodopsin
C;Keywords: G protein-coupled receptor

Query Match 42.1%; Score 780.5; DB 2; Length 378;
Best Local Similarity 42.9%; Pred. No. 3.9e-60;
Matches 153; Conservative 77; Mismatches 104; Indels 23; Gaps 6;

QY	1	MADDYGSESTSMEDYVNFNFDYCEKNVVRQFASHFLPPLYWLWVIVGALGNSLVLY	60
DB	28	VTDDYIGDNTT-----VDYTLFSLCSKCKDVRNFKAWFLPIMYSIIICFVGLLGNLVLT	82
QY	61	YWYCTRVTMTDMFLNLATADLLFLVTLFPWATAADONKQFOTFMCKVNSMYKMFYS	120
DB	83	YIYFKRLKTTDTYLLNLAVADILFLTLFPWAYSAKSWVGFVHFKCLFAIYKMFSS	142
QY	121	CVLLIMCISVDYRTAIAQAMRAHVRKRLLYSKMVCFTIWLAAALCIPILYSQKEE	180
DB	143	GMLLLICISIDRYVAIVQVSAHRHRAVLLISKLSGVHILATVLSIPELLYSDLQRS	202
QY	181	SG--IAICTWYPSDESTKLKSAVLTLLK-----ILGFFLPFVVMACCYTIITHTLIQAKS	235
DB	203	SSEQAMRCSLI-----TEHVEAFITIQVQAMVIGFLVPLPLAMSFCLYVIIRTLQARNF	256

Result No.	Query	Score	Match	Length	DB	ID	Description	
1	1854	100.0	357	1	KR9_HUMAN	P51686	homo sapien	
2	1642	88.6	369	1	KR9_MOUSE	Q9wut7	mus musculus	
3	780.5	42.1	378	1	KR7_HUMAN	P32248	homo sapien	
4	757.5	40.9	378	1	KR7_MOUSE	P47774	mus musculus	
5	652	35.2	350	1	KRB_BOVIN	P35350	bos taurus	
6	638.5	34.4	374	1	KR6_HUMAN	P51684	homo sapien	
7	637	34.4	350	1	KRB_HUMAN	Q9npb9	homo sapien	
8	624	33.7	342	1	KR6_MACNE	O19024	macaca neme	
9	619	33.4	342	1	KR6_CERAE	O18983	cercopithec	
10	617	33.3	343	1	KR6_MACMU	Q9xt45	macaca mula	
11	612	33.0	342	1	KR6_HUMAN	O00574	homo sapien	
12	606	32.7	352	1	KR5_CERTO	O62743	cercocebus	
13	605	32.6	367	1	KR6_MOUSE	O54689	mus musculus	
14	604	32.6	352	1	KR5_CERAE	P56493	cercopithec	
15	604	32.6	352	1	KR5_GORGO	P56439	gorilla gor	
16	603	32.5	352	1	KR5_MACMU	P79436	macaca mula	
17	602	32.5	352	1	KR5_PAPHIA	P56441	papio hamad	
18	601	32.4	352	1	KR5_PONPY	O97881	pongo pygma	
19	598	32.3	352	1	KR5_PANTR	P56440	pan troglod	
20	598	32.3	352	1	KR5_TRAPH	O97878	trachypithe	
21	598	32.3	352	1	KR5_TRAPH	O97878	trachypithe	
22	597	32.2	352	1	KR5_PYGBI	O97880	pygathrix b	
23	597	32.2	352	1	KR5_PYGNE	O97882	pygathrix n	
24	596	32.1	352	1	KR5_HUMAN	P51681	homo sapien	
25	595	32.1	352	1	KR5_HYLLE	O97883	hylobates l	
26	585.5	31.6	354	1	KR5_MOUSE	P51682	mus musculus	
27	585.5	31.6	360	1	KR4_HUMAN	P51679	homo sapien	
28	582	31.4	360	1	IL8B_HUMAN	P25025	homo sapien	
29	578	31.2	373	1	KR2_RAT	O55193	rattus norv	
30	577	31.1	360	1	KR4_MOUSE	P51680	mus musculus	
31	575	31.0	373	1	KR2_MOUSE	P51683	mus musculus	
32	573.5	30.9	354	1	KR5_RAT	O08556	rattus norv	
33	573	30.9	360	1	KR2_MACMU	O18793	macaca mula	

```

FT TRANSMEM 74 94 2 (POTENTIAL).
FT DOMAIN 95 108 EXTRACELLULAR (POTENTIAL).
FT TRANSMEM 109 130 3 (POTENTIAL).
FT DOMAIN 131 148 CYTOPLASMIC (POTENTIAL).
FT TRANSMEM 149 169 4 (POTENTIAL).
FT DOMAIN 170 198 EXTRACELLULAR (POTENTIAL).
FT TRANSMEM 199 226 5 (POTENTIAL).
FT DOMAIN 227 242 CYTOPLASMIC (POTENTIAL).
FT TRANSMEM 243 268 6 (POTENTIAL).
FT DOMAIN 269 292 EXTRACELLULAR (POTENTIAL).
FT TRANSMEM 293 310 7 (POTENTIAL).
FT DOMAIN 311 357 CYTOPLASMIC (POTENTIAL).
FT CARBOHYD 20 20 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT DISULFID 107 186 BY SIMILARITY.
SQ SEQUENCE 357 AA; 40713 MW; 96982E0B922F6B31 CRC64;

Query Match 100.0%; Score 1854; DB 1; Length 357;
Best Local Similarity 100.0%; Pred. No. 6.3e-115; Indels 0; Gaps 0;
Matches 357; Conservative 0; Mismatches 0;

1 MADDYSESTSMEDYVNFNFTDFCEKNNVRFASHPPLPLYWLVFVIGALGNSLVILV 60
|||||
1 MADDYSESTSMEDYVNFNFTDFCEKNNVRFASHPPLPLYWLVFVIGALGNSLVILV 60
Db

61 YWYCTRVKTMDFLLNLAIADLLFLVLPFWATAAADQWKFQTFMCKVVMNMYKMFYS 120
61 YWYCTRVKTMDFLLNLAIADLLFLVLPFWATAAADQWKFQTFMCKVVMNMYKMFYS 120
Db

121 CVLLIMCISVDRIYIAQAMRAHTWREKRLYSKMVCFIIVWLAALCIPILYSQIKEE 180
121 CVLLIMCISVDRIYIAQAMRAHTWREKRLYSKMVCFIIVWLAALCIPILYSQIKEE 180
Db

181 SGIAICTMVPSPDESTKLKSAVLTKVILGFFLPVVMACCYTIITHTLIQAKSSKHKA 240
181 SGIAICTMVPSPDESTKLKSAVLTKVILGFFLPVVMACCYTIITHTLIQAKSSKHKA 240
Db

241 LKVTITVTLVFLVSOPFYNCILLVQTDAYAMFISNCAVSTNIDICFOVQTIAFFHSCL 300
241 LKVTITVTLVFLVSOPFYNCILLVQTDAYAMFISNCAVSTNIDICFOVQTIAFFHSCL 300
Db

301 NPVLVYVGERFRDLVTKNLGICISQAQWVSFTRREGSKLSSMLLETTSGLSL 357
301 NPVLVYVGERFRDLVTKNLGICISQAQWVSFTRREGSKLSSMLLETTSGLSL 357
Db

RESULT 2
CKR9_MOUSE STANDARD; PRT; 369 AA.
CKR9_MOUSE
O9MUT7;
30-MAY-2000 (Rel. 39, Created)
30-MAY-2000 (Rel. 39, Last sequence update)
16-OCT-2001 (Rel. 40, Last annotation update)
C-C chemokine receptor type 9 (C-CR-9) (CC-CR-9) (CCR-9)
(Chemokine C-C receptor 10).
CCR9 OR CNKBR10.
Mus musculus (Mouse).
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
NCBI_TaxID=10090;
[1]
SEQUENCE FROM N.A.
Tissue=Thymus;
MEDLINE=95248139; PubMed=10229797;
Zaballos A., Gutierrez J., Varona R., Ardavin C., Marquez G.;
"Cutting edge: Identification of the orphan chemokine receptor GPR-9-6
as CCR9, the receptor for the chemokine TECK.";
J. Immunol. 162:5671-5675(1999).
-!- FUNCTION: RECEPTOR FOR CHEMOKINE SCYA25/TECK. SUBSEQUENTLY
TRANSDUCES A SIGNAL BY INCREASING THE INTRACELLULAR CALCIUM IONS
LEVEL.
-!- SUBCELLULAR LOCATION: Integral membrane protein.
-!- TISSUE SPECIFICITY: HIGHLY EXPRESSED IN THE THYMUS AND LOW IN

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CC LYMPH NODES AND SPLEEN.
CC -!- SIMILARITY: BELONGS TO FAMILY 1 OF G-PROTEIN COUPLED RECEPTORS.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL: AJ132336; CAB43480.1; -.
CC MGD: MGI:1341902; Cmkbr10.
CC InterPro: IPR004069; Chemokine9_receptor.
CC InterPro: IPR000276; GPCR_Rhodpsn.
CC Pfam: PF00001; 7tm.1; 1.
CC PRINTS: PR01531; CHEMOKINER9.
CC PRINTS: PR00237; GPCR_RHODOPSIN.
CC PROSITE: PS00237; G-PROTEIN_RECEPTOR_FL_1; 1.
CC PROSITE: PS00262; G-PROTEIN_RECEPTOR_FL_2; 1.
CC G-protein coupled receptor; Transmembrane; Glycoprotein.
CC FT DOMAIN 1 49 EXTRACELLULAR (POTENTIAL).
CC FT TRANSMEM 50 76 1 (POTENTIAL).
CC FT DOMAIN 77 85 CYTOPLASMIC (POTENTIAL).
CC FT TRANSMEM 86 106 2 (POTENTIAL).
CC FT DOMAIN 107 120 EXTRACELLULAR (POTENTIAL).
CC FT TRANSMEM 121 142 3 (POTENTIAL).
CC FT DOMAIN 143 160 CYTOPLASMIC (POTENTIAL).
CC FT TRANSMEM 161 181 4 (POTENTIAL).
CC FT DOMAIN 182 210 EXTRACELLULAR (POTENTIAL).
CC FT TRANSMEM 211 238 5 (POTENTIAL).
CC FT DOMAIN 239 254 CYTOPLASMIC (POTENTIAL).
CC FT TRANSMEM 255 280 6 (POTENTIAL).
CC FT DOMAIN 281 304 EXTRACELLULAR (POTENTIAL).
CC FT TRANSMEM 305 322 7 (POTENTIAL).
CC FT DOMAIN 323 369 CYTOPLASMIC (POTENTIAL).
CC FT CARBOHYD 32 32 N-LINKED (GLCNAC. . .) (POTENTIAL).
CC FT DISULFID 119 198 BY SIMILARITY.
CC SQ SEQUENCE 369 AA; 41913 MW; 6971F76F0A24B4AE CRC64;

Query Match 88.6%; Score 1642; DB 1; Length 369;
Best Local Similarity 86.6%; Pred. No. 4.9e-101;
Matches 309; Conservative 21; Mismatches 27; Indels 0; Gaps 0;

Qy 1 MADDYSESTSMEDYVNFNFTDFCEKNNVRFASHPPLPLYWLVFVIGALGNSLVILV 60
|||||
13 MFDDFSYDSTASTDDYNNLNFSSFFCKNNVRFASHPPLPLYWLVFVIGALGNSLVILV 72
Db

61 YWYCTRVKTMDFLLNLAIADLLFLVLPFWATAAADQWKFQTFMCKVVMNMYKMFYS 120
|||||
73. YWYCTRVKTMDFLLNLAIADLLFLVLPFWATAAADQWKFQTFMCKVVMNMYKMFYS 132
Qy 121 CVLLIMCISVDRIYIAQAMRAHTWREKRLYSKMVCFIIVWLAALCIPILYSQIKEE 180
|||||
133 CVLLIMCISVDRIYIAQAMRAHTWREKRLYSKMVCFIIVWLAALCIPILYSQIKEE 192
Db

181 SGIAICTMVPSPDESTKLKSAVLTKVILGFFLPVVMACCYTIITHTLIQAKSSKHKA 240
|||||
193 SGIAICTMVPSPDESTKLKSAVLTKVILGFFLPVVMACCYTIITHTLIQAKSSKHKA 252
Qy 241 LKVTITVTLVFLVSOPFYNCILLVQTDAYAMFISNCAVSTNIDICFOVQTIAFFHSCL 300
|||||
253 LKVTITVTLVFLVSOPFYNCILLVQTDAYAMFISNCAVSTNIDICFOVQTIAFFHSCL 312
Db

301 NPVLVYVGERFRDLVTKNLGICISQAQWVSFTRREGSKLSSMLLETTSGLSL 357
|||||
313 NPVLVYVGERFRDLVTKNLGICISQAQWVSFTRREGSKLSSMLLETTSGLSL 369
Db

RESULT 3
CKR7_HUMAN STANDARD; PRT; 378 AA.
ID CKR7_HUMAN

```

GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: October 1, 2002, 06:28:56 ; Search time 28.28 Seconds
(without alignments)
2183.847 Million cell updates/sec

Title: US-09-522-752-2
Perfect score: 1854
Sequence: 1 MADDYGSSTSMEDYVNFN.....EGSLKLSMILETTSGALS 357

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 562222 seqs, 172994929 residues

Total number of hits satisfying chosen parameters: 562222

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : SPTRMBL19:**
1: sp_archaea:*
2: sp_bacteria:*
3: sp_fungi:*
4: sp_human:*
5: sp_invertebrate:*
6: sp_mammal:*
7: sp_mhc:*
8: sp_organelle:*
9: sp_phase:*
10: sp_plant:*
11: sp_rodent:*
12: sp_virus:*
13: sp_vertebrate:*
14: sp_unclassified:*
15: sp_rvirus:*
16: sp_bacteriap:*
17: sp_archaeap:*

pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	1854	100.0	369	4 Q9UQ06	Q9uqg6 homo sapien
2	776.5	41.9	368	13 O42444	O42444 oncorhynch
3	671.5	36.2	351	11 Q9EQ16	Q9eq16 mus musculu
4	665.5	35.9	351	11 Q9ERH5	Q9erh5 mus musculu
5	645	34.8	350	11 Q924I3	Q924i3 mus musculu
6	621	33.5	343	6 Q9N020	Q9n0z0 cercocebus
7	616	33.2	342	6 Q9TV16	Q9tv16 pan troglod
8	616	33.2	352	6 Q9TV44	Q9tv44 cercoptithc
9	614	33.1	367	11 Q9RLV0	Q9rlv0 mus musculu
10	613	33.1	343	6 Q9BDS6	Q9bds6 macaca fasc.
11	612	33.0	342	4 Q9HCAS	Q9hcas homo sapien
12	612	33.0	352	6 Q9XT76	Q9xt76 cercoptithc
13	612	33.0	352	6 Q95ND1	Q95nd1 mandrillus
14	611	33.0	352	6 Q9BGN6	Q9bgn6 cercoptithc
15	608	32.8	352	6 Q95ND2	Q95nd2 mandrillus
16	607	32.7	352	6 Q9TV49	Q9tv49 cercocebus

ALIGNMENTS

RESULT 1

Q9UQ06 ID Q9UQ06 PRELIMINARY; PRT; 369 AA.
AC Q9UQ06
DT 01-MAY-2000 (TREMBLrel. 13, Created)
DT 01-MAY-2000 (TREMBLrel. 13, Last sequence update)
DT 01-OCT-2001 (TREMBLrel. 18, Last annotation update)
DE CHEMOKINE RECEPTOR CCR9 (CC CHEMOKINE RECEPTOR 9A).
GN CCR9.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP MEDLINE-99248139; PubMed-10229797;
RX Zaballos A., Gutierrez J., Varona R., Ardavin C., Marquez G.;
RT "Identification of the orphan chemokine receptor GPR-9-6 as CCR9, the
receptor for the chemokine TECK";
RL J. Immunol. 162:5671-5675(1999).
RN [2]
RP SEQUENCE FROM N.A.
RA Yu C.-R., Peden K.W.C., Farber J.M.;
RT "CCR9A and CCR9B, Two Receptors for the Chemokine CCL25 (TECK/Ckbeta-15).";
RL Submitted (APR-1999) to the EMBL/GenBank/DBJ databases.
DR EMBL; AJ132337; CAB3477.1; -
DR EMBL; AF145439; AAF66699.1; -
DR InterPro; IPR004069; Chemokine9_receptor.
DR Pfam; PF000276; GPCR_Rhodpsn.
DR PRINTS; PR01531; CHEMOKINER9.
DR PRINTS; PR02237; GPCR_RHODPSN.
DR PROSITE; PS00237; G_PROTEIN_RECFI_1; UNKNOWN_1.
DR PROSITE; PS0262; G_PROTEIN_RECFI_2; 1.
KW Receptor.
SQ SEQUENCE 369 AA; 42015 MW; F27CEA0CFB6B44C CRC64;

17 606 32.7 339 6 Q9TUQ6
18 606 32.7 352 6 Q95NE1
19 605 32.6 352 6 Q77776
20 604 32.6 339 6 Q9TQ07
21 604 32.6 339 6 Q9TSN2
22 604 32.6 352 6 Q97975
23 604 32.6 352 6 Q9TSK1
24 604 32.6 352 6 Q9TV42
25 604 32.6 352 6 Q9XS99
26 604 32.6 352 6 Q95NE8
27 604 32.6 352 6 Q95ND0
28 603 32.5 352 6 Q9XT12
29 603 32.5 352 6 Q9BGN5
30 602 32.5 352 6 Q9TV50
31 601 32.4 339 6 Q9TUW4
32 601 32.4 339 6 Q9TUS6
33 601 32.4 339 6 Q9TUS5
34 601 32.4 339 6 Q9TUR4
35 601 32.4 352 6 Q95NC1
36 600 32.4 339 6 Q9TOX3
37 600 32.4 339 6 Q9TOX2
38 600 32.4 339 6 Q9TQ05
39 600 32.4 339 6 Q9TQ04
40 600 32.4 339 6 Q9TOR2
41 600 32.4 339 6 Q9TQ05
42 600 32.4 339 6 Q9TUT6
43 600 32.4 352 6 Q9TOX0
44 600 32.4 352 6 Q95NC5
45 600 32.4 352 6 Q95NC0

Q9tuq6 erythrocebu
Q95ne1 cercocebus
Q77776 cercocebus
Q9tq07 cercoptithc
Q9tsn2 macaca fasc
Q97975 macaca arct
Q9tsk1 cercoptithc
Q9tv42 cercoptithc
Q9xs99 gorilla gor
Q95ne8 cercoptithc
Q95nd0 erythrocebu
Q9xt12 cercoptithc
Q9bgn5 cercoptithc
Q9tv50 pan troglod
Q9tuw4 pan troglod
Q9tus6 papio cynoc
Q9tus5 papio cynoc
Q9tur4 mandrillus
Q95nc1 theropithc
Q9tx3 mandrillus
Q9tx2 erythrocebu
Q9tq05 cercoptithc
Q9tq04 cercoptithc
Q9tq2 gorilla gor
Q9tu5 pan troglod
Q9tut6 macaca neme
Q9tox0 cercoptithc
Q95nc5 hylobates s
Q95nc0 hylobates m

```
Query Match 100.0%; Score 1854; DB 4; Length 369;
Best Local Similarity 100.0%; Pred. No. 1e-164;
Matches 357; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MADDYGSSTSSMEDYVNFNFTDFYCEKNNVROFASHPPLYLWLVFIVGALGNSLVTLV 60
D 13 MADDYGSSTSSMEDYVNFNFTDFYCEKNNVROFASHPPLYLWLVFIVGALGNSLVTLV 72
QY 61 YWYCTRVKTMDFMELLNLAIAIDLFLVTLPEWATAAADONKQFQFMCKVNVSMYKMFYS 120
D 73 YWYCTRVKTMDFMELLNLAIAIDLFLVTLPEWATAAADONKQFQFMCKVNVSMYKMFYS 132
QY 121 CVLLIMCISVDYRIAQAAMRAHTRKRLLYSKMVCFTIWLAAALCIPILYSQIKEE 180
D 133 CVLLIMCISVDYRIAQAAMRAHTRKRLLYSKMVCFTIWLAAALCIPILYSQIKEE 192
QY 181 SGIACTMVYPSDESTKLKSAVLTKLVILGFLPFLVFMVMACTYIIHTLQAKSSKHKA 240
D 193 SGIACTMVYPSDESTKLKSAVLTKLVILGFLPFLVFMVMACTYIIHTLQAKSSKHKA 252
QY 241 LKVTITVTLTVFLVSOFPYNCILLVQTDAYAMFISNCAVSTNIDICFQVOTQIAFFHSCL 300
D 253 LKVTITVTLTVFLVSOFPYNCILLVQTDAYAMFISNCAVSTNIDICFQVOTQIAFFHSCL 312
QY 301 NPVLYVFGVGRFRDLVKTLNKGICISOAQWVSFTRREGSKLSMLETTSGLSL 357
D 313 NPVLYVFGVGRFRDLVKTLNKGICISOAQWVSFTRREGSKLSMLETTSGLSL 369

RESULT 2
O42444 ID O42444 PRELIMINARY; PRT; 368 AA.
AC O42444;
DT 01-JAN-1998 (TrEMBLrel. 05, Created)
DT 01-JAN-1998 (TrEMBLrel. 05, Last sequence update)
DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
DE CHEMOKINE RECEPTOR.
OS Oncorhynchus mykiss (Rainbow trout) (Salmo gairdneri).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Euteleostei;
OC Protacanthopterygii; Salmoniformes; Salmonidae; Oncorhynchus.
OX NCBI_TaxID=8022;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=99260342; PubMed=10331499;
RA Daniels G.D., Zou J., Charlemagne J., Partula S., Cunningham C.,
RA Secombes C.J.;
RT "Cloning of two chemokine receptor homologs (CXC-R4 and CC-R7) in
RT rainbow trout Oncorhynchus mykiss.";
DR J. Leukoc. Biol. 65:684-690(1999).
DR EMBL; AJ003159; CAA05917.1;
DR InterPro; IPR000276; GPCR_Rhodpsn.
DR Pfam; PF00001; 7tm.1.1.
DR PRINTS; PR00237; GPCRHOOPS.
DR PROSITE; PS00237; G_PROTEIN_RECEP_FL1; UNKNOWN_1.
DR PROSITE; PS0262; G_PROTEIN_RECEP_FL2; 1.
KW Receptor.
SQ SEQUENCE 368 AA; 41523 MW; BE28E2D4C47E821A CRC64;

Query Match 41.9%; Score 776.5; DB 13; Length 368;
Best Local Similarity 43.6%; Pred. No. 2.4e-64;
Matches 157; Conservative 73; Mismatches 115; Indels 15; Gaps 6;

QY 4 DYGSSTSSMEDYVNFNFTDFYCEKNNVROFASHPPLYLWLVFIVGALGNSLVILVY- 61
D 18 DYDSFTPTVGEDVD-----NFMCKSAVAFRCQYEPPLYSIVILGGLNLTVMWYL 73
QY 62 WYCTRVKTMDFMELLNLAIAIDLFLVTLPEWATAAADONKQFQFMCKVNVSMYKMFYS 121
D 74 HFQRLKTMTDIYLLNLAIVADFLGLTLPWAVEANQGSFGLCKVTSFAFYKINFSS 133
QY 122 VLLIMCISVDYRIAQAAMRAHTRKRLLYSKMVCFTIWLAAALCIPILYSQIKEES 181
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Db 134 MLLTCTISVDYRVVIVQTTMAQNSKQRLSCSEKVCACVWLLAVLLALPEFMFANVKELD 193
QY 182 GIATCTWYVPSDESTKLKSAVLTKLVILGFLPFLVFMVMACTYIIHTLQAKSSKHKA 241
D 194 GQFCTWYVNSNRTKIVVGLQICMGFCPLPLLVWVFCYAGIIRILLKTRSTQKHKA 253
QY 242 KVTITVTLTVFLVSOFPYNCILLVQTDAYAMFISNCAVSTNIDICFQVOTQIAFFHSCL 301
D 254 RVLVWVAVFVLSQPLNSVLVMEATQAANSTQDCSAAKRFNVSVQLKSLAYTHACLN 313
QY 302 PVLVYVFGVGRFRDLVKTLNKGICISOAQWVSFTRREGSKLSMLET-TSGALS 357
D 314 PFLYVFGVGRFRDLKLLRIYHC-----WPAKGLKRIQGGPGRSSVMSDITDQALS 368

RESULT 3
Q9EQ16 ID Q9EQ16 PRELIMINARY; PRT; 351 AA.
AC Q9EQ16;
DT 01-MAR-2001 (TrEMBLrel. 16, Created)
DT 01-MAR-2001 (TrEMBLrel. 16, Last sequence update)
DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
DE CHEMOKINE RECEPTOR CXCR6.
GN CXCR6.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN=C57BL/6;
RX MEDLINE=21177382; PubMed=11017100;
RA Matloubian M., David A., Engel S., Ryan J.E., Cyster J.G.;
RT "A transmembrane CXC chemokine is a ligand for HIV-coreceptor Bonzo.";
RL Nat. Immunol. 1:298-304(2000).
DR EMBL; AF301018; AAG34367.1;
DR MGI; MGI:1934582; CXCR6.
DR InterPro; IPR000276; GPCR_Rhodpsn.
DR Pfam; PF00001; 7tm.1.1.
DR PRINTS; PR00237; GPCRHOOPS.
DR PROSITE; PS00237; G_PROTEIN_RECEP_FL1; UNKNOWN_1.
DR PROSITE; PS0262; G_PROTEIN_RECEP_FL2; 1.
KW Receptor.
SQ SEQUENCE 351 AA; 40468 MW; 5658788372B4C65A CRC64;

Query Match 36.2%; Score 671.5; DB 11; Length 351;
Best Local Similarity 40.2%; Pred. No. 1.4e-54;
Matches 137; Conservative 77; Mismatches 102; Indels 25; Gaps 8;

QY 20 NTFDYCEKNNVROFASHPPLYLWLVFIVGALGNSLVILVWYCTRVKTMDFMELLNLA 79
D 24 NSSDSENKRFLEKFEVFLPCVLYVWVFGGLGNSLVLIYFYQKRLTDFVFLNLP 83
QY 80 IADLLFLVTLPEWATAAADONKQFQFMCKVNVSMYKMFYSVLLIMCISVDYRIAQA 139
D 84 LADLVFVCTLPFWAYACTYEWVFGTVMCKTLRGMYTMNFVYVSMLTLCITVDFRIVVQA 143
QY 140 MRATWREKRLLYSKMVCFTIWLAAALCIPILYSQIKEESGIACTMVYPSDESTKLK 199
D 144 TKAFNRQAKWKGQVCLLIWVYVSLVSLPQIIYGHVQIDKL-ICQ--YHSEE---IS 197
QY 200 SAVTLKVLGFLPFLVFMVMACTYIIHTLQAKSSKHKAKTIVTLVFLVLSQFPYN 259
D 198 TWLVIVQNTLGFLPLTLMLTILCSGIKTLHARNFOKHSKLIIFLVAVFLTQTFN 257
QY 260 CILLVQTI--DAYAMFISNCAVSTNIDICFQVOTQIAFFHSCLNPVLYVFGVGRFRDLV 317
D 258 LAMLIQSTSEYVTI-----TSFYAIVTAIYAFRACLNPVLYAFVGLKFRKNV 309
QY 318 KTLKNLGICIS----QAQWVSFTRREGSKL--SSMLETTS 352
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